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## CURRENT STATUS OF ALL CLAIMS

- 1. (Currently amended) A method of determining amino acid sequence of a polypeptide, comprising:
- (a) constructing a graph from mass spectra of two or more differentially labeled polypeptides, said graph comprising a node with mass m, number of labels n, intensity i, and mass differential of labels d;
- (b) creating a node corresponding to a paired signal having masses of about m and about m+nd, [[and]]
- (c) adding a labeled weighted directed edge to said graph between any two nodes corresponding to a mass of an amino acid, said labeled weighted directed edge combining properties of said paired signals, and
- (d) assigning a satisfying amino acid to two or more of said labeled weighted directed edges, thereby determining said amino acid sequence.
- 2. (Currently amended) The method of claim 1, further comprising:
- [[(a)]] (e) creating a source node with total mass M, total number of labels N and fixed intensity  $I_{\rm s}$ ; and
- [[(b)]]  $\underline{(f)}$  creating a terminus node with mass 0, minimum number of labels  $n_0$ , and fixed intensity  $I_t$ ;
- 3. (Original) The method of claim 2, further comprising selecting a path from the source node to the terminus node.

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4. (Original) The method of claim 3, further comprising computing a priority score for each path through the graph.

- 5. (Original) The method of claim 1, wherein said differential label marks an internal amino acid residue.
- 6. (Original) The method of claim 1, wherein said differential label marks a terminal amino acid residue.
- 7. (Original) The method of claim 1, wherein said differential label marks a terminal and an internal amino acid residue.
- 8. (Original) The method of claim 1, wherein said differentially labeled polypeptides further comprise stable isotopic labels.
- 9. (Original) The method of claim 1, wherein said differentially labeled polypeptides further comprise heavy and light labeled isotopes selected from the group consisting of hydrogen, carbon, oxygen, nitrogen, sulfur and selenium.
- 10. (Original) The method of claim 1, wherein said differentially labeled polypeptides further comprise an unlabeled polypeptide and a labeled polypeptide.
- 11. (Original) The method of claim 1, wherein said polypeptide is labeled in vivo or in vitro.

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- 12. (Original) The method of claim 1, wherein said mass spectra are obtained from a mass spectrometry database.
- 13. (Original) The method of claim 1, wherein said mass spectra are of low resolution.
- 14. (Original) The method of claim 1, further comprising masses of amino acid post-translational modifications.
- 15. (Original) The method of claim 1, further comprising adding complement node with mass M-m, and a number of labels N-n+n0.
- 16. (Original) The method of claim 1, further comprising including multiple amino acid edges between nodes, said multiple amino acid edges characterizing a degenerate amino acid residue in said polypeptide sequence.
- 17. (Original) The method of claim 1, wherein steps a-c are repeated one or more times.
- 18. (Original) The method of claim 1, wherein steps a-c are performed by an automated process.

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19. (Original) A method of determining an amino acid sequence of a polypeptide, comprising:

- (a) differentially labeling two or more polypeptide mixtures, and
- (b) determining an amino acid sequence of a polypeptide within said mixture using the method of claim 1.
- 20. (Original) The method of claim 19, wherein said differential label marks an internal amino acid residue.
- 21. (Original) The method of claim 19, wherein said differential label marks a terminal amino acid residue.
- 22. (Original) The method of claim 19, wherein said differential label marks a terminal and an internal amino acid residue.
- 23. (Original) The method of claim 19, wherein said differentially labeled polypeptides further comprise stable isotopic labels.
- 24. (Original) The method of claim 19, wherein said differentially labeled polypeptides further comprise heavy and light labeled isotopes selected from the group consisting of hydrogen, carbon, oxygen, nitrogen, sulfur and selenium.
- 25. (Original) The method of claim 19, wherein said differentially labeled polypeptides further comprise an unlabeled polypeptide and a labeled polypeptide.

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- 26. (Original) The method of claim 19, wherein said polypeptide is labeled in vivo or in vitro.
- 27. (Original) The method of claim 19, wherein said mass spectra are obtained from a mass spectrometry database.
- 28. (Original) The method of claim 19, wherein said mass spectra are of low resolution.
- 29. (Original) The method of claim 19, further comprising separating components of said mixture.
- 30. (Withdrawn) A method of determining an amino acid sequence of a parent polypeptide, comprising:
- (a) obtaining mass spectra of two or more differentially labeled polypeptide fragments of a parent polypeptide;
- (b) assigning a mass and a weighting characteristic to two or more paired signals having a difference in mass corresponding to an integer value of said differential label, said weighting characteristic combining properties of each signal within said paired signals;
- (c) selecting from said mass spectra a paired signal having said assigned mass and a weighting characteristic distinguishable from non-peptide signals, said assigned mass indicating the mass of a polypeptide fragment within said spectra;
- (d) determining the difference in mass of said polypeptide fragments;
- (e) assigning said mass differences a satisfying amino acid name, and

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(f) orienting said assigned amino acid names.

- 31. (Withdrawn) The method of claim 30, wherein said differential label marks an internal amino acid residue.
- 32. (Withdrawn) The method of claim 30, wherein said differential label marks a terminal amino acid residue.
- 33. (Withdrawn) The method of claim 30, wherein said differential label marks a terminal and an internal amino acid residue.
- 34. (Withdrawn) The method of claim 30, wherein said differentially labeled polypeptides further comprise stable isotopic labels.
- 35. (Withdrawn) The method of claim 30, wherein said differentially labeled polypeptides further comprise heavy and light labeled isotopes selected from the group consisting of hydrogen, carbon, oxygen, nitrogen, sulfur and selenium.
- 36. (Withdrawn) The method of claim 30, wherein said differentially labeled polypeptides further comprise an unlabeled polypeptide and a labeled polypeptide.
- 37. (Withdrawn) The method of claim 30, wherein said parent polypeptide is labeled in vivo or in vitro.

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38. (Withdrawn) The method of claim 30, wherein said mass spectra are obtained from a mass spectrometry database.

- 39. (Withdrawn) The method of claim 30, wherein said mass spectra are of low resolution.
- 40. (Withdrawn) A method of determining an amino acid sequence of a parent polypeptide, comprising:
- (a) obtaining a mass spectra of two differentially labeled polypeptide fragments of said parent polypeptide, said differential label marking a terminal residue and at least one internal amino acid residue;
- (b) identifying a paired signal from said mass spectra corresponding to an internal amino acid residue, said paired amino acid signal having a difference in mass corresponding to said differential label;
- (c) identifying a paired signal from said mass spectra corresponding to said terminal residue, said paired amino acid signal having a difference in mass corresponding to said differential label:
- (d) determining the difference in mass of polypeptide fragments corresponding to said identified paired signals;
- (e) assigning said mass differences a satisfying amino acid name, and
  - (f) orienting said assigned amino acid names.
- 41. (Withdrawn) The method of claim 40, wherein said differential label marks two or more internal amino acid residues.

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- (Withdrawn) The method of claim 40, wherein said 42. differential label marks two terminal amino acid residues.
- 43. (Withdrawn) The method of claim 40, wherein said differential label marks a terminal and two or more internal amino acid residues.
- (Withdrawn) The method of claim 40, wherein said 44. differentially labeled polypeptides further comprise a stable isotopic label.
- (Withdrawn) The method of claim 40, wherein said differentially labeled polypeptides further comprise heavy and light labeled isotopes selected from the group consisting of hydrogen, carbon, oxygen, nitrogen, sulfur and selenium.
- The method of claim 40, wherein said 46. (Withdrawn) differentially labeled polypeptides further comprise an unlabeled polypeptide and a labeled polypeptide.
- The method of claim 40, wherein said (Withdrawn) parent polypeptide is labeled in vivo or in vitro.
- 48. (Withdrawn) The method of claim 40, wherein said mass spectra are obtained from a mass spectrometry database.
- 49. (Withdrawn) The method of claim 40, wherein said mass spectra are of low resolution.

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50. (Withdrawn) The method of claim 40, further comprising identifying a paired signal corresponding to a different internal residue having an integer difference in mass corresponding to said differential label.

- 51. (Withdrawn) The method of claim 40, further comprising identifying a paired signal corresponding to two or more internal amino acid residues having the same integer difference in mass.
- 52. (Withdrawn) The method of claim 40, wherein said step of orienting said assigned names further comprises assigning a weighted value to said paired amino acid signals.
- 53. (Withdrawn) The method of claim 40, wherein said terminal residue comprises the lowest integer difference in mass.
- 54. (Withdrawn) The method of claim 40 wherein said terminal residue is a carboxyl terminus.
- 55. (Withdrawn) The method of claim 40, wherein said terminal residue is an amino terminus.